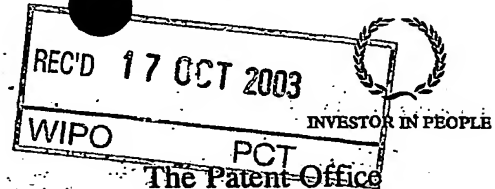


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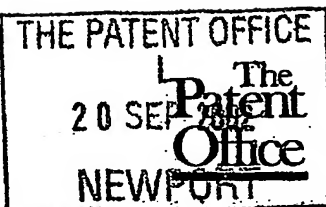
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100839

2. Patent application number

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3. Full name, address and postcode of the or of each applicant (underline all surnames)

AstraZeneca AB
S-151 85 Sodertalje
Sweden

Patents ADP number (if you know it)

If the applicant is a corporate body, give the country/state of its incorporation

Sweden

7822448003

4. Title of the invention

NOVEL COMPOUND

5. Name of your agent (if you have one)

Neil Godfrey Alasdair Phillips

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

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Claim(s)

3

Abstract

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Statement of inventorship and right to grant of a patent (Patents Form 7/77)

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NOVEL COMPOUND

The present invention relates to a thiazolopyrimidinone compound, processes and intermediates used in its preparation, pharmaceutical compositions containing it and its use in therapy.

Chemokines play an important role in immune and inflammatory responses in various diseases and disorders, including asthma and allergic diseases, as well as autoimmune pathologies such as rheumatoid arthritis and atherosclerosis. These small secreted molecules are a growing superfamily of 8-14 kDa proteins characterised by a conserved four cysteine motif. At the present time, the chemokine superfamily comprises three groups exhibiting characteristic structural motifs, the Cys-X-Cys (C-X-C), Cys-Cys (C-C) and Cys-X₃-Cys (C-X₃-C) families. The C-X-C and C-C families have sequence similarity and are distinguished from one another on the basis of a single amino acid insertion between the NH-proximal pair of cysteine residues. The C-X₃-C family is distinguished from the other two families on the basis of having a triple amino acid insertion between the NH-proximal pair of cysteine residues.

The C-X-C chemokines include several potent chemoattractants and activators of neutrophils such as interleukin-8 (IL-8) and neutrophil-activating peptide 2 (NAP-2).

The C-C chemokines include potent chemoattractants of monocytes and lymphocytes but not neutrophils. Examples include human monocyte chemotactic proteins 1-3 (MCP-1, MCP-2 and MCP-3), RANTES (Regulated on Activation, Normal T Expressed and Secreted), eotaxin and the macrophage inflammatory proteins 1 α and 1 β (MIP-1 α and MIP-1 β).

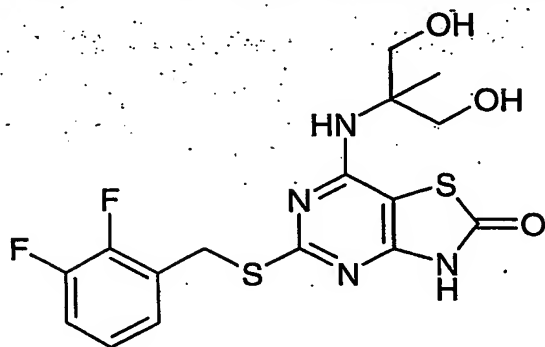
The C-X₃-C chemokine (also known as fractalkine) is a potent chemo attractant and activator of microglia in the central nervous system (CNS) as well as of monocytes, T cells, NK cells and mast cells.

Studies have demonstrated that the actions of the chemokines are mediated by subfamilies of G protein-coupled receptors, among which are the receptors designated CCR1, CCR2, CCR2A, CCR2B, CCR3, CCR4, CCR5, CCR6, CCR7, CCR8, CCR9, CCR10 and CCR11 (for the C-C family); CXCR1, CXCR2, CXCR3, CXCR4 and CXCR5 (for the C-X-C family) and CX₃CR1 for the C-X₃-C family. These receptors represent good targets for drug development since agents which modulate these receptors would be useful in the treatment of disorders and diseases such as those mentioned above.

WO-01/25242 discloses a series of thiazolopyrimidinone compounds useful as CXCR2 antagonists. A compound within the scope of WO-01/25242, but not specifically disclosed therein, has now surprisingly been found to have an improved pharmacological profile when compared with the structurally most similar compounds from WO-01/25242 i.e.

5 Examples 4 and 7.

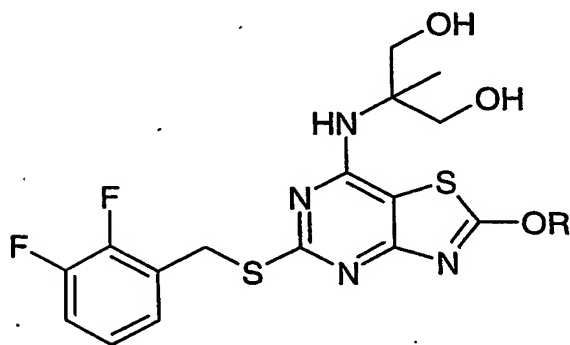
The present invention therefore provides a compound of formula (I) and pharmaceutically acceptable salts or solvates thereof:



(I)

10 The compound of formula (I) is capable of existing in tautomeric form. Tautomers and mixtures thereof also form an aspect of the present invention.

According to the invention there is also provided a process for the preparation of compound (I) which comprises reaction of a compound of formula (II):



15 (II)

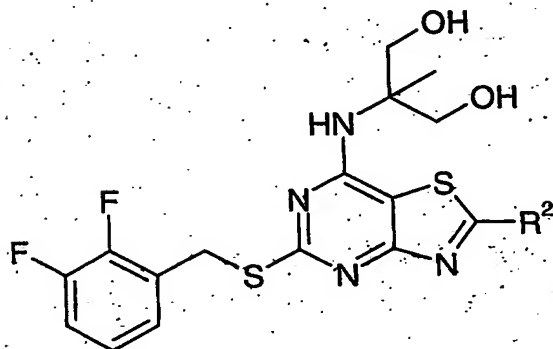
where R is C₁₋₆ alkyl with an acid,

and optionally thereafter: forming a pharmaceutically acceptable salt.

Preferably R is ethyl or methyl, more preferably methyl. Preferably the reaction is
20 carried out using dioxan and HCl. Preferably the compounds of the invention are prepared according to the procedures exemplified herein.

The compound (II) can be prepared from the corresponding compound of formula

(III):

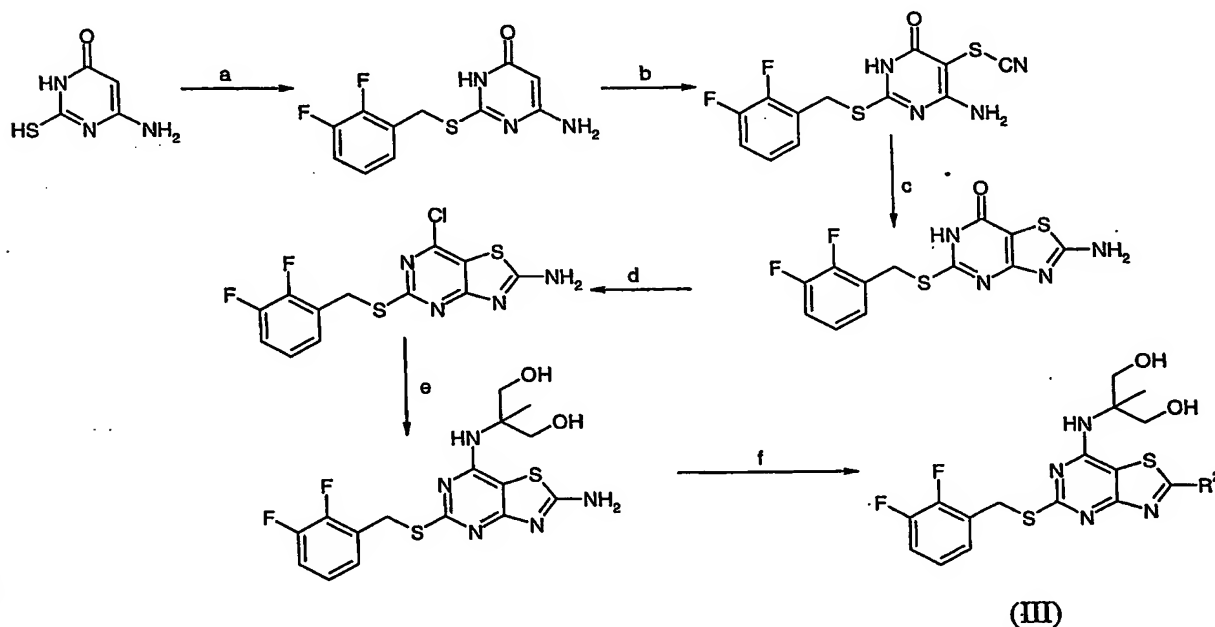


(III)

5 where R² is halogen by treating with a compound ROH in the presence of a base.

Preferably the compound of formula (III) is treated with sodium methoxide. Preferably R² is chloro.

Compounds of formula (III) can be prepared using the sequence below:



Suitable reagents for steps a to f will be known to those skilled in the art. Preferably steps a to f are carried out as exemplified herein.

15 The compound of formula (II) is itself believed to be novel and forms a further aspect of the invention.

It will be appreciated by those skilled in the art that in the processes of the present invention certain functional groups such as hydroxyl or amino groups in the starting reagents

or intermediate compound may need to be protected by protecting groups. Thus, the preparation of the compound of formula (I) may involve, at an appropriate stage, the removal of one or more protecting groups. The protection and deprotection of functional groups is fully described in 'Protective Groups in Organic Chemistry', edited by J. W. F. McOmie,

- 5 Plenum Press (1973), and 'Protective Groups in Organic Synthesis', 2nd edition, T. W. Greene & P. G. M. Wuts, Wiley-Interscience (1991).

The compound of formula (I) above may be converted to a pharmaceutically acceptable salt or solvate thereof, preferably a basic addition salt such as sodium, potassium, calcium, aluminium, lithium, magnesium, zinc, benzathine, chlorprocaine, choline,

- 10 diethanolamine, ethanolamine, ethyldiamine, meglumine, tromethamine or procaine, or an acid addition salt such as a hydrochloride, hydrobromide, phosphate, acetate, fumarate, maleate, tartrate, citrate, oxalate, methanesulphonate or *p*-toluenesulphonate.

The compound of formula (I) has activity as a pharmaceutical, in particular as a modulator of chemokine receptor (especially CXCR2) activity, and may be used in the

15 treatment (therapeutic or prophylactic) of conditions/diseases in human and non-human animals which are exacerbated or caused by excessive or unregulated production of chemokines. Examples of such conditions/diseases include:

- (1) **(the respiratory tract)** obstructive airways diseases including chronic obstructive pulmonary disease (COPD); asthma, such as bronchial, allergic,
- 20 intrinsic, extrinsic and dust asthma, particularly chronic or inveterate asthma (e.g. late asthma and airways hyper-responsiveness); bronchitis; acute, allergic, atrophic rhinitis and chronic rhinitis including rhinitis caseosa, hypertrophic rhinitis, rhinitis purulenta, rhinitis sicca and rhinitis medicamentosa; membranous rhinitis including croupous, fibrinous and pseudomembranous rhinitis and
- 25 scrofulous rhinitis; seasonal rhinitis including rhinitis nervosa (hay fever) and vasomotor rhinitis; sarcoidosis, farmer's lung and related diseases, fibroid lung and idiopathic interstitial pneumonia;
- (2) **(bone and joints)** rheumatoid arthritis, seronegative spondyloarthropathies (including ankylosing spondylitis, psoriatic arthritis and Reiter's disease),
- 30 Behcet's disease, Sjogren's syndrome and systemic sclerosis;
- (3) **(skin)** psoriasis, atopic dermatitis, contact dermatitis and other eczematous dermatides, seborrhoetic dermatitis, Lichen planus, Pemphigus, bullous Pemphigus, Epidermolysis bullosa, urticaria, angiodermas, vasculitides,

erythemas, cutaneous eosinophilias, uveitis, Alopecia areata and vernal conjunctivitis;

- 5 (4) **(gastrointestinal tract)** Coeliac disease, proctitis, eosinophilic gastro-enteritis, mastocytosis, Crohn's disease, ulcerative colitis, food-related allergies which have effects remote from the gut, e.g., migraine, rhinitis and eczema;

- 10 (5) **(central and peripheral nervous system)** Neurodegenerative diseases and dementia disorders, e.g. Alzheimer's disease, amyotrophic lateral sclerosis and other motor neuron diseases, Creutzfeldt-Jacob's disease and other prion diseases, HIV encephalopathy (AIDS dementia complex), Huntington's disease, frontotemporal dementia, Lewy body dementia and vascular dementia; polyneuropathies, e.g. Guillain-Barré syndrome, chronic inflammatory demyelinating polyradiculoneuropathy, multifocal motor neuropathy, plexopathies; CNS demyelination, e.g. multiple sclerosis, acute disseminated/haemorrhagic encephalomyelitis, and subacute sclerosing
- 15 panencephalitis; neuromuscular disorders, e.g. myasthenia gravis and Lambert-Eaton syndrome; spinal disorders, e.g. tropical spastic paraparesis, and stiff-man syndrome; paraneoplastic syndromes, e.g. cerebellar degeneration and encephalomyelitis; CNS trauma; migraine; and stroke.

- 20 (6) **(other tissues and systemic disease)** Atherosclerosis, Acquired Immunodeficiency Syndrome (AIDS), lupus erythematosus, systemic lupus, erythematosus, Hashimoto's thyroiditis, type I diabetes, nephrotic syndrome, eosinophilia fascitis, hyper IgE syndrome, lepromatous leprosy, and idiopathic thrombocytopenia purpura; post-operative adhesions, and sepsis.

- 25 (7) Stroke, subarachnoid haemorrhage, re-perfusion injury in the heart, brain, peripheral limbs and other organs.

- (8) **(allograft rejection)** acute and chronic following, for example, transplantation of kidney, heart, liver, lung, bone marrow, skin and cornea; and chronic graft versus host disease;

- 30 (9) Cancers, especially non-small cell lung cancer (NSCLC), malignant melanoma, prostate cancer and squamous sarcoma, and tumour metastasis;

- (10) Diseases in which angiogenesis is associated with raised CXCR2 chemokine levels (e.g. NSCLC, diabetic retinopathy).

- (11) Cystic fibrosis

(12) Burn wounds & chronic skin ulcers

(13) Reproductive Diseases (e.g. Disorders of ovulation, menstruation and implantation, Pre-term labour, Endometriosis)

Thus, the present invention provides a compound of formula (I), or a
5 pharmaceutically-acceptable salt or solvate thereof, as hereinbefore defined for use in therapy.

Preferably the compound of the invention is used to treat diseases in which the chemokine receptor belongs to the CXC chemokine receptor subfamily, more preferably the target chemokine receptor is the CXCR2 receptor,

Particular conditions which can be treated with the compound of the invention are
10 rheumatoid arthritis, diseases in which angiogenesis is associated with raised CXCR2 chemokine levels, and COPD. It is preferred that the compound of the invention is used to treat rheumatoid arthritis and respiratory disease.

As a further aspect of the present invention, the compound of formula (I) may have utility as an antagonist of the CX3CR1 receptor. Such a compound is expected to be
15 particularly useful in the treatment of disorders within the central and peripheral nervous system and other conditions characterized by an activation of microglia and/or infiltration of leukocytes (e.g. stroke/ischemia and head trauma).

In a further aspect, the present invention provides the use of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as hereinbefore defined in the
20 manufacture of a medicament for use in therapy.

In a still further aspect, the present invention provides the use of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as hereinbefore defined in the manufacture of a medicament for the treatment of human diseases or conditions in which modulation of chemokine receptor activity is beneficial.

25 In the context of the present specification, the term "therapy" also includes "prophylaxis" unless there are specific indications to the contrary. The terms "therapeutic" and "therapeutically" should be construed accordingly.

The invention still further provides a method of treating a chemokine mediated disease wherein the chemokine binds to a chemokine (especially CXCR2) receptor, which comprises
30 administering to a patient a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as hereinbefore defined.

The invention also provides a method of treating an inflammatory disease, especially rheumatoid arthritis, COPD, respiratory disease and psoriasis, in a patient suffering from, or

at risk of, said disease, which comprises administering to the patient a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as hereinbefore defined.

For the above-mentioned therapeutic uses the dosage administered will, of course, vary with the compound employed, the mode of administration, the treatment desired and the disorder indicated.

The compound of formula (I) and pharmaceutically acceptable salts and solvates thereof may be used on their own but will generally be administered in the form of a pharmaceutical composition in which the formula (I) compound/salt/solvate (active ingredient) is in association with a pharmaceutically acceptable adjuvant, diluent or carrier. Depending on the mode of administration, the pharmaceutical composition will preferably comprise from 0.05 to 99 %w (per cent by weight), more preferably from 0.05 to 80 %w, still more preferably from 0.10 to 70 %w, and even more preferably from 0.10 to 50 %w, of active ingredient, all percentages by weight being based on total composition.

The present invention also provides a pharmaceutical composition comprising a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as hereinbefore defined, in association with a pharmaceutically acceptable adjuvant, diluent or carrier.

The invention further provides a process for the preparation of a pharmaceutical composition of the invention which comprises mixing a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as hereinbefore defined, with a pharmaceutically acceptable adjuvant, diluent or carrier.

The pharmaceutical compositions may be administered topically (e.g. to the lung and/or airways or to the skin) in the form of solutions, suspensions, heptafluoroalkane aerosols and dry powder formulations; or systemically, e.g. by oral administration in the form of tablets, capsules, syrups, powders or granules, or by parenteral administration in the form of solutions or suspensions, or by subcutaneous administration or by rectal administration in the form of suppositories or transdermally. Preferably the compound of the invention is administered orally.

The invention will now be further illustrated by reference to the following example. Nuclear Magnetic Resonance (NMR) spectra were measured on a Varian Unity Inova 300 or 400 MHz spectrometer and the Mass Spectrometry (MS) spectra measured on a Finnigan Mat SSQ7000 or Micromass Platform spectrometer. Where necessary, the reactions were

performed under an inert atmosphere of either nitrogen or argon. Chromatography was generally performed using Matrex Silica 60[®] (35-70 micron) or Prolabo Silica gel 60[®] (35-70 micron) suitable for flash silica gel chromatography. High pressure liquid chromatography purification was performed using either a Waters Micromass LCZ with a Waters 600 pump controller, Waters 2487 detector and Gilson FC024 fraction collector or a Waters Delta Prep 4000. The abbreviations m.p. and DMSO used in the examples stand for melting point and dimethyl sulphoxide respectively.

Example 1

5-[[[(2,3-Difluorophenyl)methyl]thio]-7-[[2-hydroxy-1-(hydroxymethyl) 1

10 methylethyl]amino]-thiazolo[4,5-*d*]pyrimidin-2(3*H*)-one

a) 6-Amino-2-[[[(2,3-difluorophenyl)methyl]thio]- 4(3*H*)-pyrimidinone

4-Amino-6-hydroxy-2-mercaptopyrimidine monohydrate (7.1g) was added portion wise to a stirred suspension of 60% sodium hydride (2.4g) in dry *N,N*-dimethylformamide (70ml).

After 1 hour a solution of 2,3-Difluorobenzyl bromide (10g) in dry *N,N*-dimethylformamide (10ml) was added. Stirred over weekend at room temperature. Poured on to ice/water and the precipitate was collected by filtration to give 9.6g of product. 81% yield.

MS (APCI) (+ve) 270 (M+H, 94%)

b) 6-Amino-2-[[[(2,3-difluorophenyl)methyl]thio]-1,6-dihydro-6-oxo-5-pyrimidinyl ester thiocyanic acid

20 The product from step (a) (28g) and potassium thiocyanate (40.5g) in *N,N*-dimethylformamide (583ml) were heated together at 65°C. Pyridine (14.5ml) was added and the solution cooled to 5°C. Bromine (5.0ml) was added slowly and the reaction mixture stirred for 2 hours at 5-10°C. The reaction mixture was poured onto ice water (4200ml), stirred for 1 hour and the solid was collected by filtration, washed with water and ether, to
25 give 24g of product. 70% yield.

MS (APCI) (+ve) 327 (M+H)

c) 2-Amino-5-[[[(2,3-difluorophenyl)methyl]thio]-thiazolo[4,5-*d*]pyrimidin-7(6*H*)-one

A mixture of the product from step (b) (12.1g), *N,N*-dimethylformamide (70ml) and water (20ml) was heated to 120°C for 24 hours. A colourless solid precipitated from the solution,
30 which was allowed to cool, and the solid collected by filtration to give 8.3g of product. 70% yield.

MS (APCI) (+ve) 327 (M+H)

d) 7-Chloro-5-[[2,3-difluorophenyl)methyl]thio]-thiazolo[4,5-*d*]pyrimidin-2-amine

The product of step (c) (10.0g) was suspended in phosphoryl chloride (55ml). *N,N*-dimethylaniline (5.5ml) added slowly and reaction mixture heated at reflux for 2 hours. Allowed to cool, then poured on to ice with vigorous stirring; temperature was not allowed to go above 45°C (ice added). After approximately 20 minutes the temperature stabilized at 30°C. The solid that formed was collected by filtration and washed with water. Purified by column chromatography (EtOAc to 5% MeOH in EtOAc) to give 3.34g of product. 31% yield.

MS: APCI (+ve) 345 (M+H)

10 e) 2-[[2-Amino-5-[[2,3-difluorophenyl)methyl]thio]thiazolo[4,5-*d*]pyrimidin-7-yl]amino]-2-methyl-1,3-propanediol

The product from step (d) (1.5g) was suspended in NMP (10ml), then Hunigs base (1.5ml) and 2-Amino-2-methylpropanediol (1.37g) added. Reaction mixture heated to 110°C under N₂ for 4 hrs. A further aliquot of 2-Amino-2-methylpropanediol (0.685g) was added and
15 mixture heated at 110°C for 5 hrs. Mixture poured in to water (400ml) and solid was collected by filtration. Purified by column chromatography (EtOAc: Methanol (95:5)) to give 0.756g of product. 42% yield.

MS: APCI (+ve) 414 (M+H)

20 f) 2-[[2-Chloro-5-[[2,3-difluorophenyl)methyl]thio]thiazolo[4,5-*d*]pyrimidin-7-yl]amino]-2-methyl-1,3-propanediol

The product from step (e) (0.485g) was suspended in conc.HCl (18ml) which was then cooled to 15°C. A mixture of water (15ml) and acetonitrile (25ml) added to give a solution. Cooled to 5°C and a solution of sodium nitrite (0.162g) in water (1ml) added drop-wise. Stirred at 5°C for several hours then allowed to warm overnight. Solution cooled to -10°C and
25 neutralized with ammonia, then concentrated in vacuo. The yellow precipitate was collected by filtration and washed with water. Dried in vacuo to give 0.339g of product. 67% yield.

MS: APCI (+ve) 433 (M+H)

g) 2-[[5-[[2,3-Difluorophenyl)methyl]thio]-2-methoxythiazolo[4,5-*d*]pyrimidin-7-yl]amino]-2-methyl-1,3-propanediol

30 The product from step (f) (0.339g) was suspended in methanol (32ml). Potassium hydroxide (0.088g) added and mixture stirred at 50°C for 20 minutes. Neutralised with 2N HCl and solvents removed in vacuo to give an orange residue. Water added to remove inorganics and the yellow solid was collected by filtration to give 0.3g of desired product. 90% yield.

MS: APCI (+ve) 429 (M+H)

h) 5-[[2,3-Difluorophenyl)methyl]thio]-7-[[2-hydroxy-1-(hydroxymethyl)-1-methylethyl]amino]-thiazolo[4,5-*d*]pyrimidin-2(3*H*)-one

5 The product from step (g) (0.3g) was suspended in a mixture of dioxan (50ml) and conc.HCl (1ml). Water (1ml) was added and the resultant solution heated at 60°C for 12 hours.

Allowed to stand over weekend. Solvents removed in vacuo and residue taken up in water.

Yellow precipitate collected by filtration and washed with water. Purified using prep. Hplc Acetonitrile:0.1% ammonium acetate (90:10 to 95:5) over 25 minutes to give 0.063g of the

10 desired product. 22% yield.

MS: APCI (+ve) 415 (M+H)

¹H NMR: δ (DMSO) 1.25 (3H, s), 3.54-3.66 (4H, m), 4.39 (2H, s), 4.65-4.69 (2H, t), 6.34 (1H, s), 7.12-7.20 (1H, m), 7.29-7.41 (2H, m), 12.43 (1H, s).

mp 230-233 °C

15 Example 2

5-[[2,3-Difluorophenyl)methyl]thio]-7-[[2-hydroxy-1-(hydroxymethyl)-1-methylethyl]amino]-thiazolo[4,5-*d*]pyrimidin-2(3*H*)-one, monosodium salt.

The product from Example 1, step (h) (0.87g) was suspended in water (80ml), 1.0M sodium hydroxide (3.0ml) added, then methanol (15ml), and the mixture heated on a steam

20 bath. When dissolution was almost complete, the mixture was filtered hot and the filtrate chilled overnight to give a fluffy white precipitate after filtration.

This was dried in a vacuum oven overnight at 50°C. (0.60g).

MS: APCI (+ve) 415 (M+H)

¹H NMR: δ (DMSO) 1.23 (3H, s), 3.47-3.58 (4H, m), 4.37 (2H, s), 4.94 (2H, t), 5.29 (1H, s),

25 7.14 (1H, m), 7.31 (1H, m), 7.35 (1H, m).

mp 238 °C (dec.)

Pharmacological Data

Ligand Binding Assay

30 [¹²⁵I]IL-8 (human, recombinant) was purchased from Amersham, U.K. with a specific activity of 2,000Ci/mmol. All other chemicals were of analytical grade. High levels of hrCXCR2 were expressed in HEK 293 cells (human embryo kidney 293 cells ECACC No. 85120602) (Lee *et al.* (1992) *J. Biol. Chem.* **267** pp16283-16291). hrCXCR2 cDNA was amplified and

- cloned from human neutrophil mRNA. The DNA was cloned into PCRScript (Stratagene) and clones were identified using DNA. The coding sequence was sub-cloned into the eukaryotic expression vector RcCMV (Invitrogen). Plasmid DNA was prepared using Quiagen Megaprep 2500 and transfected into HEK 293 cells using Lipofectamine reagent (Gibco BRL). Cells of the highest expressing clone were harvested in phosphate-buffered saline containing 0.2%(w/v) ethylenediaminetetraacetic acid (EDTA) and centrifuged (200g, 5min.). The cell pellet was resuspended in ice-cold homogenisation buffer [10mM HEPES (pH 7.4), 1mM dithiothreitol, 1mM EDTA and a panel of protease inhibitors (1mM phenyl methyl sulphonyl fluoride, 2µg/ml soybean trypsin inhibitor, 3mM benzamidine, 0.5µg/ml leupeptin and 100µg/ml bacitracin)] and the cells left to swell for 10 minutes. The cell preparation was disrupted using a hand held glass mortar/PTFE pestle homogeniser and cell membranes harvested by centrifugation (45 minutes, 100,000g, 4°C). The membrane preparation was stored at -70°C in homogenisation buffer supplemented with Tyrode's salt solution (137mM NaCl, 2.7mM KCl, 0.4mM NaH₂PO₄), 0.1%(w/v) gelatin and 10%(v/v) glycerol.
- All assays were performed in a 96-well MultiScreen 0.45µm filtration plates (Millipore, U.K.). Each assay contained ~50pM [¹²⁵I]IL-8 and membranes (equivalent to ~200,000 cells) in assay buffer [Tyrode's salt solution supplemented with 10mM HEPES (pH 7.4), 1.8mM CaCl₂, 1mM MgCl₂, 0.125mg/ml bacitracin and 0.1%(w/v) gelatin]. In addition, a compound of formula (I) according to the Examples was pre-dissolved in DMSO and added to reach a final concentration of 1%(v/v) DMSO. The assay was initiated with the addition of membranes and after 1.5 hours at room temperature the membranes were harvested by filtration using a Millipore MultiScreen vacuum manifold and washed twice with assay buffer (without bacitracin). The backing plate was removed from the MultiScreen plate assembly, the filters dried at room temperature, punched out and then counted on a Cobra γ-counter.
- The compound of formula (I) has an IC₅₀ value of less than (<) 10µM.

Intracellular Calcium Mobilisation Assay

- Human neutrophils were prepared from EDTA-treated peripheral blood, as previously described (Baly *et al.* (1997) *Methods in Enzymology* 287 pp70-72), in storage buffer [Tyrode's salt solution (137mM NaCl, 2.7mM KCl, 0.4mM NaH₂PO₄) supplemented with 5.7mM glucose and 10mM HEPES (pH 7.4)].

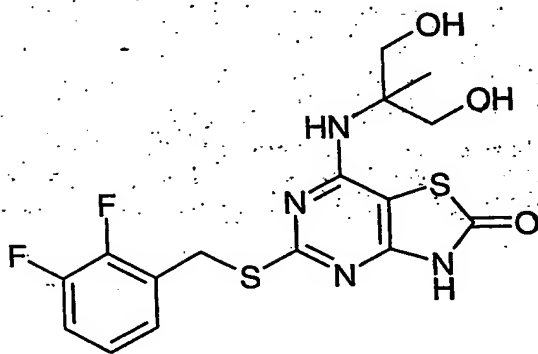
The chemokine GRO α (human, recombinant) was purchased from R&D Systems (Abingdon, U.K.). All other chemicals were of analytical grade. Changes in intracellular free calcium were measured fluorometrically by loading neutrophils with the calcium sensitive fluorescent dye, fluo-3, as described previously (Merritt *et al.* (1990) *Biochem. J.* 269, pp513-519). Cells
5 were loaded for 1 hour at 37°C in loading buffer (storage buffer with 0.1%(w/v) gelatin) containing 5 μ M fluo-3 AM ester, washed with loading buffer and then resuspended in Tyrode's salt solution supplemented with 5.7mM glucose, 0.1%(w/v) bovine serum albumin (BSA), 1.8mM CaCl₂ and 1mM MgCl₂. The cells were pipetted into black walled, clear bottom, 96 well micro plates (Costar, Boston, U.S.A.) and centrifuged (200g, 5 minutes, room
10 temperature).

A compound of formula (I) according to the Examples was pre-dissolved in DMSO and added to a final concentration of 0.1%(v/v) DMSO. Assays were initiated by the addition of an A₅₀ concentration of GRO α and the transient increase in fluo-3 fluorescence (λ_{Ex} = 490nm and λ_{Em} = 520nm) monitored using a FLIPR (Fluorometric Imaging Plate Reader, Molecular
15 Devices, Sunnyvale, U.S.A.).

The compound of formula (I) was tested and found to be an antagonist of the CXCR2 receptor in human neutrophils.

CLAIMS

1. A compound of formula (I) and pharmaceutically acceptable salts or solvates thereof:

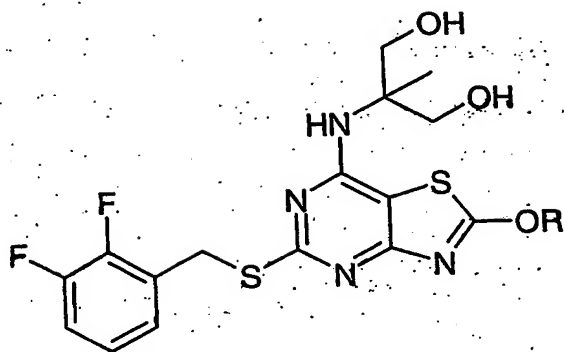


5

(I)

2. A pharmaceutical composition comprising a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as claimed in claim 1 in association with a pharmaceutically acceptable adjuvant, diluent or carrier.
3. A process for the preparation of a pharmaceutical composition as claimed in claim 2 which comprises mixing a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as claimed in claim 1 with a pharmaceutically acceptable adjuvant, diluent or carrier.
4. A compound of formula (I), or a pharmaceutically-acceptable salt or solvate thereof, as claimed in claim 1 for use in therapy.
5. Use of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as claimed in claim 1 in the manufacture of a medicament for use in therapy.
6. A method of treating a chemokine mediated disease wherein the chemokine binds to one or more chemokine receptors, which comprises administering to a patient a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as claimed in claim 1.

7. A method according to claim 6 in which the chemokine receptor belongs to the CXC chemokine receptor subfamily.
8. A method according to claim 6 or 7 in which the chemokine receptor is the CXCR2
5 receptor.
9. A method of treating an inflammatory disease in a patient suffering from, or at risk of, said disease, which comprises administering to the patient a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as
10 claimed in claim 1.
10. A method according to claim 9, wherein the disease is psoriasis.
11. A method according to claim 9, wherein the disease is rheumatoid arthritis.
15
12. A method of treating a disease in which angiogenesis is associated with raised CXCR2 chemokine levels, or at risk of, said disease, which comprises administering to the patient a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as claimed in claim 1.
20
13. A method according to claim 9, wherein the disease is severe asthma.
14. A method according to claim 9, wherein the disease is inflammatory bowel disease.
- 25 15. A method according to claim 9, wherein the disease is COPD.
16. A process for the preparation of a compound of formula (I) which comprises reaction of a compound of formula (II):



(II)

- 5 where R is C₁₋₆ alkyl with an acid,
and optionally thereafter: forming a pharmaceutically acceptable salt.

ABSTRACT

Title: Novel Compound

The invention provides a thiazolone compound for use in therapy.

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